

Welcome, ctrptrialsubmitter3 | Sign Out Role: trial submitter | Write Mode: On

Study Source:\* National \* Lead Organization Trial Identifier:\* QA - Team Exercitationem in de Other Trial Identifier: Protocol ID Type Protocol ID

Select a Protocol ID Ty T + Add 
 Protocol ID Type
 Protocol ID

 Other Identifier
 654654645
 Phase:\* I Clinical Research Category:\* Interventional Primary Purpose:\* Treatment Accrual Disease Terminology:\* SDC Lead Organization:\* Harborview Medical Center Q Search Organizations Principal Investigator:\* Lee, James Q Search Persons Sponsor:\* Harborview Medical Center Data Table 4 Funding Source:\* Q Search Organizations Memorial Hospital Colorado Springs Program code: e5345355 NIH Grant Information (for NIH funded Trials) Is this trial funded by an NCI grant?\*  $\odot$  Yes  $\ \ \, \odot$  No Status Date Status Comment

Select a Trial Status

Select a Trial Status Why Study Stopped + Add Status Date Status Comment

Exercitationem in deserunt rerum voluptatem. Provident, of u.o. adjoinant, of u.o. adjoinant Does this trial have an associated ○ Yes ● No IND/IDE?:\* The information in this section is REQUIRED to enable "Upload from NCI CTRP" in ClinicalTrials.gov: Responsible Party: Sponsor-Investigator Investigator:\* Lee, James Q Search Persons Investigator Title:\* Principal Investigator Investigator Affiliation:\* Harborview Medical Center Country Organization

-Select a Country- Y Select an Organization- Y + Add FDA Regulated Intervention Indicator: 

No Pes N/A Section 801 Indicator: 

No Se Data Monitoring Committee Appointed ◎ No ◎ Yes ● N/A Indicator: To ensure successful registration, upload a Protocol document and an IRB Approval document. If the Protocol document does not include the Informed Consent and/or participating sites, upload the Informed Consent document and a list of participating sites separately. You can use the Participating Sites template to submit your list of participating sites. Protocol Document:\* Choose File No file chosen 1Sample.pdf IRB Approval:\* Choose File No file chosen List of Participating Sites: Choose File No file chosen Informed Consent Document: Choose File No file chosen + Add More ... Created By: ctrptrialsubmitter3 (05-Oct-2016 9:38) \* Reset Submit Save as Draft



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View Trial

Trial Identifiers:

Lead Organization Trial Identifier: QA - Team Exercitationem in de

NCI ID: NCI-2016-01600

 Other Trial Identifier:
 Protocol ID Type
 Protocol ID

 Other
 654654645

Official Title:

QA - Team Title - Exercitationem in deserunt rerum voluptatem. Provident, sit, quo adipisioing laboriosam, quibusdam molestae quis aliquam aliquin et minima eum quo excepturi repellendus. Maxime molestae molestae sess, exercitationem ea sed sit dolor sequi quibusdam qui sint, qua erperlendent, prodent, voluptate repellendus. Eligend facilis maiores quia eus optio, veniam, eco, et quis reprehendent assumenda eligendi culps, perferendis ratione nilini eld omnis culpa provident, aliqua. Ext., que modi rure voluptate fugiat maiores recusandae. Alias est praesentum harum officia salpia

Pilot? Yes

Clinical Research Category: Interventional Primary Purpose: Treatment

Secondary Purpose: Ancillary-Correlative Accrual Disease Terminology: SDC

Lead Organization/Principal Investigator:

Lead Organization: Harborview Medical Center

Principal Investigator: Lee, James

Sponsor: Harborview Medical Center

Data Table 4 Information:

Study Source: National

Data Table 4 Funding Source: Memorial Hospital Colorado Springs

Program Code: e5345355

Current Trials Status: In Review Current Trials Status Date: 05-Oct-2016

Trial Dates:

Trial Start Date: 06-Jul-2016 Actual Primary Completion Date: 06-Jul-2018 Anticipated Completion Date: 06-Jul-2020 Anticipated

Regulatory Information:

Responsible Party: Sponsor-Investigator

Investigator: Lee, James

Investigator Title: Principal Investigator

Investigator Affiliation: Harborview Medical Center

FDA Regulated Intervention Indicator: N/A

Section 801 Indicator: N/A

Data Monitoring Committee Appointed N/A Indicator:

Protocol Document 1Sample.pdf IRB Approval 1Sample.pdf

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